PTC/SB/08x (08-03)
Approved for use through 07/31/2006 ONE 0651-0201
U.S. Patent and Trademark Office, U.S. DEPARTMENT OF COMMERCE
to a collection of information unfers it contains a valid OMS control number. Under the Paperwork Reduction Act of 1995, no persons are required to

	Application Number		10570480	
	Filing Date		2006-03-01	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)	First Named Inventor BARI		DACHENKO	
	Art Unit			
	Examiner Name			
	Attorney Docket Numb	er	5616VB-1	
	II S PATENTS		Remove	

			U.S.PATENTS			Remove				
Examiner Initial*	Cite No	Patent Number	Kind Code ¹			Name of Patentee or Applicant of cited Document		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		
	1	6337919		2002-0	8-01	Duntun				
If you wis	h to a	dd additional U.S. Pater	nt citatio	n inform	ation pl	ease click the	Add button.	_	Add	
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	tion	n Name of Patentee or Applicant of cited Document		Releva	Columns,Lines wher ant Passages or Rele s Appear	
	1									
If you wis	h to a	dd additional U.S. Publi		p				d button		
				FOREIG	SN PAT	ENT DOCUM	IENTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	e or	Pages,Columns,Lines where Relevant Passages or Relevan Figures Appear	74
	1	2097519	RU			1977-11-27	Bardachenko Vital Feodosievich	y		
	2	2126168	RU			1999-02-10	Varshavsky Zinov Matveevich	,		
	3	2360677	GB			2001-09-26	Pilot Systems (Lor Ltd.	idon)		

	Application Number		10570480	
INFORMATION DIGGS COURT	Filing Date		2006-03-01	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	First Named Inventor BAR		DACHENKO	
(Not for submission under 37 CFR 1.99)	Art Unit			
(NOTION SAUSINGSSON GINGER OF OTTE 1.504)	Examiner Name			
	Attorney Docket Numb	er	5616VB-1	

	4	1013	0019	DE		2003-01-09	SCM Microsystems GmbH				
If you wis	If you wish to add additional Foreign Patent Document citation information please click the Add button Add										
	NON-PATENT LITERATURE DOCUMENTS Remove										
Examiner Initials*	Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of th							T6			
	1 PCT International Search Report dated January 20, 2005, for PCT Application No. PCTRIA2004000069							Z			
If you wish to add additional non-patent literature document citation information please click the Add button Add											
EXAMINER SIGNATURE											
Examiner	Signa	ture					Date Considered				

See Kint Codes of USPTO Patent Documents at InventUSPTO.GDV or MPEP 901.04. Enter office that issued the document, by the Nov-letter option Standard 513.) For Laplanese patent focuments, the inclusion of the parent of the Improver protects the sent number the patent document. A focument by the appropriate symbols as enducated on the document under WIPO Standard 51.16 if possible. Spapicant is to place a check mark here if English language translation is attached.

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

Attorney Docket Number

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s):

CERTIFICATION STATEMENT

5616VR-1

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filling of the information disclosure statement. See 37 CFR 197(eVT).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1.59(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.39(c) and the statement. See 37 CFR 1.39(c) and the statement. See 37 CFR 1.39(c) and the statement. See

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

□ None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Bradley M. Knepper/	Date (YYYY-MM-DD)	2006-10-03
Name/Print	Bradley M. Knepper	Registration Number	44189

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 and 3T CFR.

1.14. This collection is estimated to take I hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operatment of Commence, P. O. Box 1430, Alexandriu, V.S. 2213.1450, DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.2.231.1450.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uting an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2014 and 2016. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the control of t
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.